## Congress of the United States

Washington, DC 20515

May 22, 2025

The Honorable Tom Cole U.S. House of Representatives H-307 The Capitol Washington, DC 20515

The Honorable Ken Calvert U.S. House of Representatives H-405 The Capitol Washington, DC 20515

The Honorable Robert Aderholt U.S. House of Representatives 2358-B Rayburn House Office Building Washington, DC 20515 The Honorable Rosa DeLauro U.S. House of Representatives 2413 Rayburn House Office Building Washington, DC 20515

The Honorable Betty McCollum U.S. House of Representatives 2426 Rayburn House Office Building Washington, DC 20515

Dear Chairman Cole, Ranking Member DeLauro, Chairman Calvert, Ranking Member McCollum, and Chairman Aderholt:

We thank you for your inclusion of funding related to lupus research in Fiscal Year 2025 Appropriations legislation. In support of the Congressional Lupus Caucus and on behalf of the hundreds of thousands of Americans living with lupus, we are writing to express our support for critically important lupus programs in the Fiscal Year 2026 Department of Defense and Labor, Health and Human Services and Education appropriations bills.

## We respectfully request the following in the Department of Defense appropriations bill:

<u>\$15 million</u> for the Congressionally Directed Medical Research Program's (CDMRP) Lupus Research Program. Research has found that those who serve in the military may be at greater risk of developing lupus.

**Program**. Research has found that those who serve in the military may be at greater risk of developing lupus. Post-traumatic stress disorder, which is experienced by 20 to 30 percent of service members, has been associated with a twofold increase in risk for autoimmune diseases like lupus, and there are likely other risk factors associated with military service. Additionally, it is imperative to advance understanding of biological factors of lupus, which will improve treatment and potentially cut medical costs of patients in the military. Over a 10-year period, nearly 68,000 lupus service members and beneficiaries were seen for lupus. Robust funding for the LRP will support the development of innovative, high-impact research to advance understanding of lupus and improve outcomes for our active-duty service members, veterans, their families, and other Americans with lupus.

We respectfully request the following in the Labor, Health and Human Services, and Education, and Related Agencies appropriations bill:

**<u>\$20 million</u>** for the Centers for Disease Control and Prevention (CDC), Chronic Prevention and Health Promotion, National Lupus Patient Registry. Since 2003, Congress has provided funding for the National Lupus Registry Program, and the program has made important strides in lupus research and education. For

example, the program funded research to help determine incidence and prevalence of the disease in the United States, including how the disease impacts different populations across the country, and identified risk factors to developing the disease and those that contribute to worse health outcomes.

The program has also developed tools and resources and launched initiatives to help improve care for people with lupus. For example, funding provided by Congress through the Registry led to the development of the first and only online self-management program and efforts to reduce the time it takes to diagnose lupus, which can be as long as six years. This program has played a critical role in improving the health of Americans living with lupus and empowering people with the tools they need to live with lupus and manage their own care.

## <u>\$3 million</u> for the Office of the HHS Secretary, Office of Minority Health (OMH), National Lupus

**Training, outreach and Clinical Trial Program.** Funding provided by Congress has enabled the lupus community to identify barriers to participation in clinical trials, a significant problem in ensuring the safety and efficacy of treatments for diseases like lupus that are more prevalent in women and in women of color. The program also has implemented action plans to increase participation in lupus clinical trials and currently is supporting multiple projects throughout the country that are actively working to ensure that trials are representative of the actual patient population, which can enable trials to more accurately assess the efficacy and safety of new treatments in everyone living with lupus, improving health outcomes and quality of life. Moreover, representative trials can ensure that medications are appropriately prescribed and covered by insurance and available to the patients who need them, reducing unnecessary medical expenses and saving lives.

Funding in FY2026 is needed to continue these programs and support our constituents whose lives have been impacted by lupus.

Thank you for your support of these initiatives.

Sincerely,

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Andrew R. Garbarino Member of Congress

Jasmine Crockett Member of Congress

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William R. Keating Member of Congress

David Scott Member of Congress

Lloyd Doggett Member of Congress

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